

2.5 Switching Patients To or From a Monoamine Oxidase Inhibitor


At least 14 days (2 weeks) should elapse between discontinuation of an MAOI and initiation of therapy with LUVOX CR Capsules. Similarly, at least 14 days (2 weeks) should be allowed after stopping LUVOX CR Capsules before starting an MAOI (4, 5.2).


2.6 Discontinuation of Treatment with LUVOX CR Capsules

Symptoms associated with discontinuation of other SSRIs or SNRIs have been reported (see **WARNINGS AND PRECAUTIONS [5.10]**). Patients should be monitored for these symptoms when discontinuing treatment. A gradual reduction in the dose rather than abrupt cessation is recommended whenever possible. If intolerable symptoms occur following a decrease in the dose or upon discontinuation of treatment, then resuming the previously prescribed dose may be considered. Subsequently, the health care provider may continue decreasing the dose but at a more gradual rate.

3 DOSAGE FORMS AND STRENGTHS

LUVOX CR Capsules are available as:

100 mg Extended-Release Capsules: a two-piece gelatin capsule (dark blue opaque cap/white opaque body) imprinted with  on one side of the cap and LCR 100 on the other side of the cap.

150 mg Extended-Release Capsules: a two-piece gelatin capsule (dark blue opaque cap/powder blue opaque body) imprinted with  on one side of the cap and LCR 150 on the other side of the cap.

4 CONTRAINDICATIONS

Coadministration of thioridazine, tizanidine, pimozide, alosetron, or ramelteon with LUVOX CR Capsules is contraindicated (see **WARNINGS AND PRECAUTIONS [5.4-5.8]**).

The use of MAOIs in combination with LUVOX CR Capsules, or within 14 days (2 weeks) of discontinuing treatment with LUVOX CR Capsules, is contraindicated. Also, LUVOX CR Capsules should not be administered within 14 days (2 weeks) after discontinuing treatment with an MAOI (see **WARNINGS AND PRECAUTIONS [5.2]**).

5 WARNINGS AND PRECAUTIONS

5.1 Clinical Worsening and Suicide Risk

Patients with major depressive disorder (MDD), both adult and pediatric, may experience worsening of their depression and/or the emergence of suicidal ideation and behavior (suicidality) or unusual changes in behavior, whether or not they are taking antidepressant medications, and this risk may persist until significant remission occurs. Suicide is a known risk of depression and certain other psychiatric disorders, and these disorders themselves are the strongest predictors of suicide. There has been a long-standing concern, however, that antidepressants may have a role in inducing worsening of depression and the emergence of suicidality in certain patients during the early phases of treatment. The pooled analyses of short-term placebo-controlled trials of antidepressant drugs (SSRIs and others) showed that these drugs increase the risk of suicidal thinking and behavior (suicidality) in children, adolescents, and young adults (ages 18-24) with major depressive disorder (MDD) and other psychiatric disorders. Short-term studies did not show an increase in the risk of suicidality with antidepressants compared to placebo in adults beyond age 24; there was a reduction with antidepressants compared to placebo in adults age 65 and older.

The pooled analyses of placebo-controlled trials in children and adolescents with MDD, obsessive compulsive disorder (OCD), or other psychiatric disorders included a total of 24 short-term trials of 9 antidepressant drugs in over 4400 patients. The pooled analyses of placebo-controlled trials in adults with MDD or other psychiatric disorders included a total of 295 short-term trials (median duration of 2 months) of 11 antidepressant drugs in over 77,000 patients. There was considerable variation in risk of suicidality among drugs, but a tendency toward an increase in the younger patients for almost all drugs studied. There were differences in absolute risk of suicidality across the different indications, with the highest incidence in MDD. The risk differences (drug vs placebo), however, were relatively stable within age strata and across indications. These risk differences (drug-placebo difference in the number of cases of suicidality per 1000 patients treated) are provided in [Table 1](#).

TABLE 1
DRUG-PLACEBO DIFFERENCES IN NUMBER
OF CASES OF SUICIDALITY PER 1000 PATIENTS TREATED

Age Range	Drug-Related Increases
<18	14 additional cases
18-24	5 additional cases
Age Range	Drug-Related Decreases
25-64	1 fewer case
≥ 65	6 fewer cases

No suicides occurred in any of the pediatric trials. There were suicides in the adult trials, but the number was not sufficient to reach any conclusion about the drug effect on suicide.

It is unknown whether the suicidality risk extends to longer-term use, i.e., beyond several months. However, there is substantial evidence from placebo-controlled maintenance trials in adults with depression that the use of antidepressants can delay the recurrence of depression.

All patients being treated with antidepressants for any indication should be monitored appropriately and observed closely for clinical worsening, suicidality, and unusual changes in behavior, especially during the initial few months of a course of drug therapy, or at times of dose changes, either increases or decreases.

The following symptoms, anxiety, agitation, panic attacks, insomnia, irritability, hostility, aggressiveness, impulsivity, akathisia (psychomotor restlessness), hypomania, and mania, have been reported in adult and pediatric patients being treated with antidepressants for major depressive disorder as well as for other indications, both psychiatric and nonpsychiatric. Although a causal link between the emergence of such symptoms and either the worsening of depression and/or the emergence of suicidal impulses has not been established, there is concern that such symptoms may represent precursors to emerging suicidality.

Consideration should be given to changing the therapeutic regimen, including possibly discontinuing the medication, in patients whose depression is persistently worse, or who are experiencing emergent suicidality or symptoms that might be precursors to worsening depression or suicidality, especially if these symptoms are severe, abrupt in onset, or were not part of the patient's presenting symptoms.

If the decision has been made to discontinue treatment, medication should be tapered, as rapidly as is feasible, but with recognition that abrupt discontinuation can be associated with certain symptoms (see *DOSAGE AND ADMINISTRATION—Discontinuation of Treatment with LUVOX CR Capsules* [5.10], for a description of the risks of discontinuation of LUVOX CR Capsules).

Families and caregivers of patients being treated with antidepressants for major depressive disorder or other indications, both psychiatric and nonpsychiatric, should be alerted about the need to monitor patients for the emergence of agitation, irritability, unusual changes in behavior, and the other symptoms described above, as well as the emergence of suicidality, and to report such symptoms immediately to health care providers. Such monitoring should include daily observation by families and caregivers. Prescriptions for LUVOX CR Capsules should be written for the smallest quantity of capsules consistent with good patient management, in order to reduce the risk of overdose.

Screening Patients for Bipolar Disorder: A major depressive episode may be the initial presentation of bipolar disorder. It is generally believed (though not established in controlled trials) that treating such an episode with an antidepressant alone may increase the likelihood of precipitation of a mixed/manic episode in patients at risk for bipolar disorder. Whether any of the symptoms described above represent such a conversion is unknown. However, prior to initiating treatment with an antidepressant, patients with depressive symptoms should be adequately screened to determine if they are at risk for bipolar disorder; such screening should include a detailed psychiatric history, including a family history of suicide, bipolar disorder, and depression. It should be noted that LUVOX CR Capsules are not approved for use in treating bipolar depression.

5.2 Potential for Monoamine Oxidase Inhibitors Interaction

In patients receiving another serotonin reuptake inhibitor drug in combination with monoamine oxidase inhibitors (MAOIs), there have been reports of serious, sometimes fatal, reactions including hyperthermia, rigidity, myoclonus, autonomic instability with possible rapid fluctuations of vital signs, and mental status changes that include extreme agitation progressing to delirium and coma. These reactions have also been reported in patients who have discontinued that drug and have been started on an MAOI. Some cases presented with features resembling a serotonin syndrome or Neuroleptic Malignant Syndrome. Therefore, LUVOX CR Capsules should not be used in combination with an MAOI, or within 14 days (2 weeks) of discontinuing treatment with an MAOI. Similarly, at least 14 days (2 weeks) should be

allowed after stopping LUVOX CR Capsules before starting an MAOI (see *DOSAGE AND ADMINISTRATION* [2.5] and *CONTRAINDICATIONS* [4]).

5.3 Serotonin Syndrome or Neuroleptic Malignant Syndrome (NMS)-like Reactions

The development of a potentially life-threatening serotonin syndrome or Neuroleptic Malignant Syndrome (NMS)-like reactions have been reported with SNRIs and SSRIs alone, including LUVOX CR treatment, but particularly with concomitant use of serotonergic drugs (including triptans), with drugs that impair metabolism of serotonin (including MAOIs), or with antipsychotics or other dopamine antagonists. Serotonin syndrome symptoms may include mental status changes (e.g., agitation, hallucinations, coma), autonomic instability (e.g., tachycardia, labile blood pressure, hyperthermia), neuromuscular aberrations (e.g., hyperreflexia, incoordination) and/or gastrointestinal symptoms (e.g., nausea, vomiting, diarrhea). Serotonin syndrome, in its most severe form, can resemble Neuroleptic Malignant Syndrome, which includes hyperthermia, muscle rigidity, autonomic instability with possible rapid fluctuation of vital signs, and mental status changes. Patients should be monitored for the emergence of serotonin syndrome or NMS-like signs and symptoms.

The concomitant use of LUVOX CR Capsules with MAOIs intended to treat depression is contraindicated.

If concomitant treatment of LUVOX CR Capsules with a 5-hydroxytryptamine receptor agonist (triptan) is clinically warranted, careful observation of the patient is advised, particularly during treatment initiation and dose increases.

The concomitant use of LUVOX CR Capsules with serotonin precursors (such as tryptophan) is not recommended.

Treatment with LUVOX CR Capsules and any concomitant serotonergic or antidopaminergic agents, including antipsychotics, should be discontinued immediately if the above reactions occur and supportive symptomatic treatment should be initiated.

5.4 Potential Thioridazine Interaction

The effect of fluvoxamine (25 mg immediate-release tablets given twice daily for one week) on thioridazine steady-state concentrations was evaluated in 10 male inpatients with schizophrenia. Concentrations of thioridazine and its two active metabolites, mesoridazine and sulforidazine, increased 3-fold following coadministration of fluvoxamine.

Thioridazine administration produces a dose-related prolongation of the QTc interval, which is associated with serious ventricular arrhythmias, such as torsades de pointes-type arrhythmias, and sudden death. It is likely that this experience underestimates the degree of risk that might occur with higher doses of thioridazine. Moreover, the effect of fluvoxamine may be even more pronounced when it is administered at higher doses.

Therefore, LUVOX CR Capsules should not be coadministered with thioridazine (see *CONTRAINDICATIONS* [4]).

5.5 Potential Tizanidine Interaction

Fluvoxamine is a potent inhibitor of CYP1A2, and tizanidine is a CYP1A2 substrate. The effect of immediate-release fluvoxamine maleate tablets (100 mg daily for four days) on the pharmacokinetics and pharmacodynamics of a single dose of tizanidine has been studied in 10 healthy male subjects. Tizanidine C_{max} was increased approximately 12-fold (range 5-fold to 32-fold), elimination half-life was increased by almost 3-fold, and AUC increased 33-fold (range 14-fold to 103-fold). The mean maximal effect on blood pressure was a 35 mm Hg decrease in systolic blood pressure, a 20 mm Hg decrease in diastolic blood pressure, and a 4 beat/min decrease in heart rate. Drowsiness was significantly increased, and performance on the psychomotor task was significantly impaired. LUVOX CR Capsules and tizanidine should not be used together (see *CONTRAINDICATIONS* [4]).

5.6 Potential Pimozide Interaction

Pimozide is metabolized by the CYP3A4 isozyme and it has been demonstrated that ketoconazole, a potent inhibitor of CYP3A4, blocks the metabolism of this drug, resulting in increased plasma concentrations of parent drug. An increased plasma concentration of pimozide causes QT prolongation and has been associated with torsades de pointes-type ventricular tachycardia, sometimes fatal. As noted below, a substantial pharmacokinetic interaction has been observed for fluvoxamine in combination with alprazolam, a drug that is known to be metabolized by the CYP3A4 isozyme. Although it has not been definitively demonstrated that fluvoxamine is a potent CYP3A4 inhibitor, it is likely to be, given the substantial interaction of fluvoxamine with alprazolam. Consequently, it is recommended that fluvoxamine not be used in combination with pimozide (see *CONTRAINDICATIONS* [4]).

5.7 Potential Alosetron Interaction

In a pharmacokinetic study, 40 healthy female subjects received fluvoxamine in escalating doses from 50 to 200 mg/day for 16 days with coadministration of alosetron 1 mg on the last day. Fluvoxamine increased mean alosetron plasma concentrations (AUC) approximately 6-fold and prolonged the half-life by approximately 3-fold. Consequently, it is recommended that LUVOX CR Capsules not be used in combination with alosetron (see **CONTRAINDICATIONS** [4] and Lotronex[®] (aloksetron) package insert).

5.8 Potential Ramelteon Interaction

When immediate-release fluvoxamine maleate tablets 100 mg twice daily were administered for 3 days prior to single-dose coadministration of ramelteon 16 mg and immediate-release fluvoxamine maleate tablets, the AUC for ramelteon increased approximately 190-fold and the C_{max} increased approximately 70-fold compared to ramelteon administered alone. Ramelteon should not be used in combination with LUVOX CR Capsules (see **CONTRAINDICATIONS** [4]).

5.9 Other Potentially Important Drug Interactions

Benzodiazepines: Benzodiazepines metabolized by hepatic oxidation (e.g., alprazolam, midazolam, triazolam, etc.) should be used with caution because the clearance of these drugs is likely to be reduced by fluvoxamine. The clearance of benzodiazepines metabolized by glucuronidation (e.g., lorazepam, oxazepam, temazepam) is unlikely to be affected by fluvoxamine.

Alprazolam - When immediate-release fluvoxamine maleate tablets (100 mg given once daily) and alprazolam (1 mg given 4 times per day) were coadministered to steady state, plasma concentrations and other pharmacokinetic parameters (AUC, C_{max} , $T_{1/2}$) of alprazolam were approximately twice those observed when alprazolam was administered alone; oral clearance was reduced by about 50%. The elevated plasma alprazolam concentrations resulted in decreased psychomotor performance and memory. This interaction, which has not been investigated using higher doses of fluvoxamine, may be more pronounced if a 300 mg daily dose is coadministered, particularly since fluvoxamine exhibits non-linear pharmacokinetics over the dosage range 100-300 mg. If alprazolam is coadministered with LUVOX CR Capsules, the initial alprazolam dosage should be at least halved and titration to the lowest effective dose is recommended. No dosage adjustment is required for LUVOX CR Capsules.

Diazepam - The coadministration of LUVOX CR Capsules and diazepam is generally not advisable. Because fluvoxamine reduces the clearance of both diazepam and its active metabolite, N-desmethyldiazepam, there is a strong likelihood of substantial accumulation of both species during chronic coadministration.

Evidence supporting the conclusion that it is inadvisable to coadminister fluvoxamine and diazepam is derived from a study in which healthy volunteers taking 150 mg/day of immediate-release fluvoxamine maleate tablets were administered a single oral dose of 10 mg of diazepam. In these subjects (N=8), the clearance of diazepam was reduced by 65% and that of N-desmethyldiazepam to a level that was too low to measure over the course of the 2-week-long study.

It is likely that this experience significantly underestimates the degree of accumulation that might occur with repeated diazepam administration. Moreover, as noted with alprazolam, the effect of fluvoxamine may even be more pronounced when it is administered at higher doses.

Accordingly, diazepam and fluvoxamine should not ordinarily be coadministered.

Clozapine: Elevated serum levels of clozapine have been reported in patients taking immediate-release fluvoxamine maleate tablets and clozapine. Since clozapine-related seizures and orthostatic hypotension appear to be dose related, the risk of these adverse reactions may be higher when fluvoxamine and clozapine are coadministered. Patients should be closely monitored when LUVOX CR Capsules and clozapine are used concurrently.

Methadone: Significantly increased methadone (plasma level:dose) ratios have been reported when immediate-release fluvoxamine maleate tablets were administered to patients receiving maintenance methadone treatment, with symptoms of opioid intoxication in one patient. Opioid withdrawal symptoms were reported following fluvoxamine maleate discontinuation in another patient.

Mexiletine: The effect of steady-state fluvoxamine (50 mg given twice daily for 7 days) on the single dose pharmacokinetics of mexiletine (200 mg) was evaluated in 6 healthy Japanese males. The clearance of mexiletine was reduced by 38% following coadministration with fluvoxamine compared to mexiletine alone. If fluvoxamine and mexiletine are coadministered, serum mexiletine levels should be monitored.

Theophylline: The effect of steady-state immediate-release fluvoxamine maleate tablets (50 mg tablets given twice daily) on the pharmacokinetics of a single dose of theophylline (375 mg as 442 mg aminophylline) was evaluated in 12 healthy non-smoking, male volunteers. The clearance of theophylline was decreased approximately 3-fold. Therefore, if theophylline is coadministered with fluvoxamine maleate, its dose should be reduced to one-third of the usual daily maintenance dose and plasma concentrations of theophylline should be monitored. No dosage adjustment is required for LUVOX CR Capsules.

Warfarin and Other Drugs That Interfere With Hemostasis (NSAIDs, Aspirin, etc.): Serotonin release by platelets plays an important role in hemostasis. Epidemiological studies of the case-control and cohort design have demonstrated an association between use of psychotropic drugs that interfere with serotonin reuptake and the occurrence of upper gastrointestinal bleeding. These studies have also shown that concurrent use of an NSAID or aspirin may potentiate this risk of bleeding. Thus, patients should be cautioned about the use of such drugs concurrently with fluvoxamine (see **WARNINGS AND PRECAUTIONS-Abnormal Bleeding [5.11]**).

Warfarin - When immediate-release fluvoxamine maleate tablets (50 mg given three times daily) were administered concomitantly with warfarin for two weeks, warfarin plasma concentrations increased by 98% and prothrombin times were prolonged. Thus patients receiving oral anticoagulants and LUVOX CR Capsules should have their prothrombin time monitored and their anticoagulant dose adjusted accordingly. No dosage adjustment is required for LUVOX CR Capsules.

5.10 Discontinuation of Treatment with LUVOX CR Capsules

During marketing of immediate-release fluvoxamine maleate tablets and other SSRIs and SNRIs (serotonin and norepinephrine reuptake inhibitors), there have been spontaneous reports of adverse reactions occurring upon discontinuation of these drugs, particularly when abrupt, including the following: dysphoric mood, irritability, agitation, dizziness, sensory disturbances (e.g., paresthesias, such as electric shock sensations), anxiety, confusion, headache, lethargy, emotional lability, insomnia, and hypomania. While these reactions are generally self-limiting, there have been reports of serious discontinuation symptoms.

Patients should be monitored for these symptoms when discontinuing treatment with LUVOX CR Capsules. A gradual reduction in the dose rather than abrupt cessation is recommended whenever possible. If intolerable symptoms occur following a decrease in the dose or upon discontinuation of treatment, then resuming the previously prescribed dose may be considered. Subsequently, the health care provider may continue decreasing the dose but at a more gradual rate (see **DOSAGE AND ADMINISTRATION [2.6]**).

5.11 Abnormal Bleeding

SSRIs and SNRIs, including LUVOX CR Capsules, may increase the risk of bleeding events. Concomitant use of aspirin, nonsteroidal anti-inflammatory drugs, warfarin, and other anticoagulants may add to this risk. Case reports and epidemiological studies (case-control and cohort design) have demonstrated an association between use of drugs that interfere with serotonin reuptake and the occurrence of gastrointestinal bleeding. Bleeding events related to SSRIs and SNRIs use have ranged from ecchymoses, hematomas, epistaxis, and petechiae to life-threatening hemorrhages.

Patients should be cautioned about the risk of bleeding associated with the concomitant use of LUVOX CR Capsules and NSAIDs, aspirin, or other drugs that affect coagulation (see **WARNINGS AND PRECAUTIONS [5.9]**).

5.12 Activation of Mania/Hypomania

During premarketing studies of immediate-release fluvoxamine maleate tablets involving primarily depressed patients, hypomania or mania occurred in approximately 1% of patients treated with fluvoxamine. In a 10-week pediatric OCD study, 2 out of 57 patients (4%) treated with fluvoxamine experienced manic reactions, compared to none of 63 placebo patients. Activation of mania/hypomania has also been reported in a small proportion of patients with major affective disorder who were treated with other marketed antidepressants. As with all antidepressants, LUVOX CR Capsules should be used cautiously in patients with a history of mania.

5.13 Seizures

During premarketing studies with immediate-release fluvoxamine maleate tablets, seizures were reported in 0.2% of fluvoxamine-treated patients. Caution is recommended when the drug is administered to patients with a history of convulsive disorders. Fluvoxamine should be avoided in patients with unstable epilepsy, and patients with controlled epilepsy should be carefully monitored. Treatment with fluvoxamine should be discontinued if seizures occur or if seizure frequency increases.

5.14 Hyponatremia

Hyponatremia may occur as a result of treatment with SSRIs and SNRIs, including LUVOX CR Capsules. In many cases, this hyponatremia appears to be the result of the syndrome of inappropriate antidiuretic hormone secretion (SIADH). Cases with serum sodium lower than 110 mmol/L have been reported. Elderly patients may be at greater risk of developing hyponatremia with SSRIs and SNRIs (see *USE IN SPECIFIC POPULATIONS, Geriatric Use [8.5]*). Also, patients taking diuretics or who are otherwise volume depleted may be at greater risk. Discontinuation of LUVOX CR Capsules should be considered in patients with symptomatic hyponatremia and appropriate medical intervention should be instituted.

Signs and symptoms of hyponatremia include headache, difficulty concentrating, memory impairment, confusion, weakness, and unsteadiness, which may lead to falls. Signs and symptoms associated with more severe and/or acute cases have included hallucination, syncope, seizure, coma, respiratory arrest, and death.

5.15 Use in Patients with Concomitant Illness

Closely monitored clinical experience with LUVOX CR Capsules in patients with concomitant systemic illness is limited. Caution is advised in administering LUVOX CR Capsules to patients with diseases or conditions that could affect hemodynamic responses or metabolism.

LUVOX CR Capsules or immediate-release fluvoxamine maleate tablets have not been evaluated or used to any appreciable extent in patients with a recent history of myocardial infarction or unstable heart disease. Patients with these diagnoses were systematically excluded from many clinical studies during premarketing testing of these products. Evaluation of the electrocardiograms for patients with depression or OCD who participated in premarketing studies revealed no differences between fluvoxamine and placebo in the emergence of clinically important ECG changes.

Patients with Hepatic Impairment - In patients with liver dysfunction, following administration of immediate-release fluvoxamine maleate tablets, fluvoxamine clearance was decreased by approximately 30%. Patients with liver dysfunction should begin with a low dose of LUVOX CR Capsules and increase it slowly with careful monitoring.

5.16 Laboratory Tests

There are no specific laboratory tests recommended.

6 ADVERSE REACTIONS

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in clinical practice.

6.1 Clinical Trial Data Sources

LUVOX CR Capsules were studied in one 12-week controlled trial in patients with OCD (N = 124; mean exposure 66.6 days) and in two 12-week controlled trials for another condition (N = 279; mean exposure 59.2 days). Patients in these trials were initiated on 100 mg/day and were titrated in 50 mg increments over the first 6 weeks to within a range of 100 mg to 300 mg/day. The reactions listed in [Table 2](#) show reactions from the two populations separately. [Table 3](#) shows reactions from the three controlled studies combined.

6.2 Adverse Reactions Observed in Controlled Trials

Adverse Reactions Associated with Discontinuation of Treatment: Of the 124 patients with OCD and 279 patients in other studies treated with LUVOX CR Capsules in controlled clinical trials, 19% and 26% discontinued treatment due to an adverse reaction. The most common reactions ($\geq 1\%$) associated with discontinuation and considered to be drug related (i.e., those reactions associated with dropout at a rate at least twice that of placebo) were *anorexia* (including, but not limited to, loss of appetite and decreased appetite) (1%), *anxiety* (3%), *asthenia* (3%), *diarrhea* (2%), *dizziness* (4%), *headache* (2%), *insomnia* (5%), *nausea* (7%), *nervousness* (1%), *somnolence* (5%), and *thinking abnormal* (1%).

Commonly Observed Adverse Reactions: LUVOX CR Capsules have been studied in one controlled trial in patients with OCD (N = 124) and two controlled trials for another condition (N = 279). In general, adverse reaction rates were similar in the two data sets as well as in a study of pediatric patients with OCD treated with immediate-release fluvoxamine maleate tablets. The most commonly observed treatment-emergent adverse reactions associated with the use of LUVOX CR Capsules and likely to be drug-related (incidence of 5% or greater and at least twice that for placebo) and derived from [Table 2](#) were: *abnormal ejaculation*, *anorexia*, *anorgasmia*, *asthenia*, *diarrhea*, *nausea*, *somnolence*, *sweating*, and *tremor*. In the one controlled trial in patients with OCD, the following additional reactions occurred at an incidence of 5% or greater and at least twice that for placebo: *anxiety*, *decreased libido*, *myalgia*, *pharyngitis*, and *vomiting*. The following additional reactions occurred in another studied population:

dyspepsia, dizziness, insomnia, and yawning. In a study evaluating immediate-release fluvoxamine maleate tablets in pediatric patients with OCD, the following additional reactions were identified using the above rule: *agitation, depression, dysmenorrhea, flatulence, hyperkinesia, and rash.*

Adverse Reactions Occurring at an Incidence of $\geq 2\%$: Table 2 enumerates adverse reactions that occurred in adults at a frequency of 2% or more, and were more frequent than in the placebo group, among patients treated with LUVOX CR Capsules in two short-term, placebo-controlled trials (12 weeks) in another population and one short-term placebo-controlled OCD trial (12 weeks) and in which patients were dosed once-a-day in a range of 100 to 300 mg/day. This table shows the percentage of patients in each group who had at least one occurrence of a reaction at some time during their treatment. Reported adverse reactions were classified using a COSTART-based Dictionary terminology.

The prescriber should be aware that these figures cannot be used to predict the incidence of side effects in the course of usual medical practice where patient characteristics and other factors may differ from those that prevailed in the clinical trials. Similarly, the cited frequencies cannot be compared with figures obtained from other clinical investigations involving different treatments, uses, and investigators. The cited figures, however, do provide the prescribing health care provider with some basis for estimating the relative contribution of drug and non-drug factors to the side-effect incidence rate in the population studied.

TABLE 2
TREATMENT-EMERGENT ADVERSE REACTION INCIDENCE RATES
BY BODY SYSTEM IN ADULT
OCD PATIENTS AND ANOTHER STUDIED POPULATION¹

BODY SYSTEM/ ADVERSE REACTION	PERCENTAGE OF PATIENTS REPORTING REACTION			
	OBSESSIVE COMPULSIVE DISORDER		OTHER STUDIED POPULATION	
	LUVOX CR N = 124	PLACEBO N = 124	LUVOX CR N = 279	PLACEBO N = 276
BODY AS A WHOLE				
Headache	32	31	35	30
Asthenia	26	8	24	10
Pain ²	10	8	—	—
Abdominal Pain	--	--	5	4
Accidental Injury	5	3	—	—
Chest Pain	--	--	3	1
Viral Infection	2	<1	—	—
CARDIOVASCULAR				
Palpitation	--	--	3	1
Vasodilatation	--	--	2	<1
Hypertension	2	<1	—	—
DIGESTIVE SYSTEM				
Nausea	34	13	39	11
Diarrhea	18	8	14	5
Anorexia ³	13	5	14	1
Dyspepsia	8	5	10	4
Constipation	4	<1	6	5
Vomiting	6	2	—	—
Tooth Disorder	2	<1	—	—
Liver Function Test Abnormal	--	--	2	<1
Gingivitis	2	0	—	—
HEMIC AND LYMPHATIC				
Ecchymosis	4	2	—	—
METABOLIC AND NUTRITIONAL DISORDERS				
Weight Loss	2	<1	—	—
MUSCULOSKELETAL				
Myalgia	5	2	—	—
NERVOUS SYSTEM				
Insomnia	35	20	32	13
Somnolence	27	11	26	9
Dizziness	12	10	15	7

BODY SYSTEM/ ADVERSE REACTION	PERCENTAGE OF PATIENTS REPORTING REACTION OBSESSIVE COMPULSIVE DISORDER		OTHER STUDIED POPULATION	
	LUVOX CR N = 124	PLACEBO N = 124	LUVOX CR N = 279	PLACEBO N = 276
Dry Mouth	10	9	11	8
Nervousness	--	--	10	9
Libido Decreased	6	2	6	4
Male	10	5	8	6
Female	4	1	4	3
Anxiety	6	2	8	5
Tremor	6	0	8	<1
Abnormal Thinking	3	<1	3	2
Abnormal Dreams	--	--	3	2
Agitation	2	<1	3	<1
Hypertonia	--	--	2	1
Apathy	3	0	–	–
Paresthesia	--	--	3	2
Neurosis	2	<1	–	–
Twitching	2	0	–	–
RESPIRATORY SYSTEM				
Pharyngitis	6	<1	–	–
Yawn	2	0	5	<1
Laryngitis	3	0	–	–
Bronchitis	--	--	2	1
Epistaxis	2	0	–	–
SKIN				
Sweating	7	<1	6	2
Acne	2	0	–	–
SPECIAL SENSES				
Taste Perversion	2	<1	2	<1
Amblyopia	2	<1	–	–
UROGENITAL				
Abnormal Ejaculation	10	0	11	2
Anorgasmia	5	0	5	1
Male	4	0	4	2
Female	5	0	5	0
Menorrhagia	3	0	–	–
Sexual Function Abnormal	2	<1	3	<1
Male	4	3	2	1
Female	0	0	3	0
Urinary Tract Infection	--	--	2	<1
Polyuria	2	<1	–	–

¹ Events for which fluvoxamine maleate incidence was equal to or less than placebo include the following for OCD patients: abdominal pain, flu syndrome, infection, palpitation, flatulence, increased appetite, weight gain, abnormal dreams, amnesia, hypertonia, nervousness, paresthesia, increased cough, dyspnea, rhinitis, and ear pain. In the other studied population the following events were seen: abdominal pain, accidental injury, back pain, flu syndrome, infection, pain, flatulence, pharyngitis, rhinitis, rash, and dysmenorrhea.

² Term includes body aches/pains, dental pain, pain from surgery, unspecified pain, and general pain secondary to injuries (sprains, fractures).

³ Includes, but is not limited to, loss of appetite and decreased appetite.

6.3 Other Adverse Reactions in OCD Pediatric Population

In pediatric patients (N=57) treated with immediate-release fluvoxamine maleate tablets, the overall profile of adverse reactions was generally similar to that seen in adult studies, as shown in Table 2. However, the following adverse reactions, not appearing in Table 2, were reported in two or more of the pediatric patients and were more

frequent with immediate-release fluvoxamine maleate tablets than with placebo: cough increase, dysmenorrhea, ecchymosis, emotional lability, epistaxis, hyperkinesia, manic reaction, rash, sinusitis, and weight decrease.

6.4 Male and Female Sexual Dysfunction with SSRIs

Although changes in sexual desire, sexual performance, and sexual satisfaction often occur as manifestations of a psychiatric disorder and with aging, they may also be a consequence of pharmacologic treatment. In particular, some evidence suggests that selective serotonin reuptake inhibitors (SSRIs) can cause such untoward sexual experiences.

Reliable estimates of the incidence and severity of untoward experiences involving sexual desire, performance, and satisfaction are difficult to obtain, however, in part because patients and health care providers may be reluctant to discuss them. Accordingly, estimates of the incidence of untoward sexual experience and performance cited in product labeling are likely to underestimate their actual incidence.

Table 3 displays the incidence of sexual side effects reported by at least 2% of patients taking LUVOX CR Capsules in placebo-controlled trials.

TABLE 3
PERCENTAGE OF PATIENTS REPORTING SEXUAL ADVERSE REACTIONS IN
PLACEBO-CONTROLLED TRIALS

	LUVOX CR N = 403	Placebo N = 400
Abnormal Ejaculation	11	2
Anorgasmia		
Male	4	1
Female	5	0
Impotence	2	2
Libido Decreased		
Male	8	5
Female	4	2
Sexual Function Abnormal		
Male	3	1
Female	2	0

Fluvoxamine treatment has been associated with several cases of priapism. In those cases with a known outcome, patients recovered without sequelae and upon discontinuation of fluvoxamine.

While it is difficult to know the precise risk of sexual dysfunction associated with the use of SSRIs, health care providers should routinely inquire about such possible side effects.

6.5 Weight and Vital Sign Changes

No statistically significant differences in weight gain or loss were found between patients treated with LUVOX CR Capsules or placebo. Comparisons of immediate-release fluvoxamine maleate tablets or LUVOX CR Capsules versus placebo groups in separate short-term trials on (1) median change from baseline on various vital signs variables and on (2) incidence of patients meeting criteria for potentially important changes from baseline on various measures of vital signs variables revealed no important differences between fluvoxamine maleate and placebo.

6.6 Laboratory Changes

Comparisons of immediate-release fluvoxamine maleate tablets or LUVOX CR Capsules versus placebo groups in separate short-term trials on (1) median change from baseline on various serum chemistry, hematology, and urinalysis variables and on (2) incidence of patients meeting criteria for potentially important changes from baseline on various serum chemistry, hematology, and urinalysis variables revealed no important differences between fluvoxamine maleate and placebo.

6.7 ECG Changes

Comparisons of immediate-release fluvoxamine maleate tablets or LUVOX CR Capsules and placebo groups in separate pools of short-term OCD and depression trials on (1) mean change from baseline on various ECG variables and on (2) incidence of patients meeting criteria for potentially important changes from baseline on various ECG variables revealed no important differences between fluvoxamine maleate and placebo.

6.8 Other Reactions Observed During the Premarketing Evaluation of Fluvoxamine

During premarketing clinical trials conducted in North America and Europe, multiple doses of LUVOX CR Capsules or immediate-release fluvoxamine maleate tablets were administered for a combined total of 3219 patient exposures in patients suffering OCD or other studied disorders. These exposures include 482 patient exposures with LUVOX CR Capsules and 2737 patient exposures with immediate-release fluvoxamine maleate tablets. Untoward reactions associated with this exposure were recorded by clinical investigators using descriptive terminology of their own choosing. Consequently, it is not possible to provide a meaningful estimate of the proportion of individuals experiencing adverse reactions without first grouping similar types of untoward reactions into a limited (i.e., reduced) number of standard reaction categories.

In the tabulations that follow, a COSTART-based Dictionary terminology has been used to classify reported adverse reactions. If the COSTART term for a reaction was so general as to be uninformative, it was replaced with a more informative term when possible. The frequencies presented, therefore, represent the proportion of the total patient exposures to multiple doses of fluvoxamine maleate who experienced a reaction of the type cited on at least one occasion while receiving fluvoxamine maleate. Reactions are further classified within body system categories and enumerated in order of decreasing frequency using the following definitions: frequent adverse reactions are defined as those occurring on one or more occasions in at least 1/100 patients; infrequent adverse reactions are those occurring between 1/100 and 1/1000 patients; and rare adverse reactions are those occurring in less than 1/1000 patients. It is important to emphasize that, although the events reported did occur during treatment with fluvoxamine maleate, a causal relationship to fluvoxamine maleate has not been established.

For LUVOX CR, all reported events are included in the list below, with the following exclusions: 1) those events already listed in [Table 2](#) or previous sections of this prescribing information; 2) those events for which there is no basis to suspect a causal relationship; and 3) events that were reported in only one patient and judged not to be potentially serious.

Body as a Whole: Infrequent: chills, malaise, photosensitivity reaction, suicide attempt.

Cardiovascular System: Infrequent: syncope.

Digestive System: Infrequent: eructation, increased salivation.

Metabolic and Nutritional Disorders: Frequent: weight gain.

Nervous System: Infrequent: confusion, incoordination, sleep disorder, suicidal tendency.

Skin and Appendages: Infrequent: eczema, urticaria.

Special Senses: Infrequent: dry eyes, photophobia, taste loss.

Urogenital System: Infrequent: vaginal hemorrhage¹.

¹Based on the number of females.

For immediate-release fluvoxamine tablets, all reported events are included in the list below, with the following exclusions: 1) those events already listed in [Table 2](#), in previous sections of this prescribing information, or in the LUVOX CR list of [Other Reactions Observed During Premarketing Evaluation](#); 2) those events for which there is no basis to suspect a causal relationship; and 3) events that were reported in only one patient and judged not to be potentially serious.

Body as a Whole: Infrequent: allergic reaction, neck pain, neck rigidity, overdose; **Rare:** sudden death.

Cardiovascular System: Frequent: hypotension; **Infrequent:** angina pectoris, bradycardia, cardiomyopathy, cardiovascular disease, cold extremities, conduction delay, myocardial infarction, pallor, pulse irregular, ST segment changes; **Rare:** AV block, cerebrovascular accident, embolus, pericarditis, phlebitis, pulmonary infarction, supraventricular extrasystoles.

Digestive System: **Frequent:** elevated liver transaminases; **Infrequent:** colitis, esophagitis, gastritis, gastroenteritis, gastrointestinal hemorrhage, gastrointestinal ulcer, glossitis, hemorrhoids, melena, rectal hemorrhage, stomatitis; **Rare:** biliary pain, cholecystitis, cholelithiasis, fecal incontinence, hematemesis, intestinal obstruction, jaundice.

Endocrine System: **Infrequent:** hypothyroidism; **Rare:** goiter.

Hemic and Lymphatic Systems: **Infrequent:** leukocytosis, lymphadenopathy, thrombocytopenia; **Rare:** leukopenia, purpura.

Metabolic and Nutritional Systems: **Frequent:** edema; **Infrequent:** dehydration, hypercholesterolemia; **Rare:** diabetes mellitus, hyperglycemia, hyperlipidemia, hypoglycemia, hypokalemia, lactate dehydrogenase increased.

Musculoskeletal System: **Infrequent:** arthralgia, arthritis, bursitis, generalized muscle spasm, myasthenia; **Rare:** myopathy.

Nervous System: **Frequent:** amnesia, apathy, hyperkinesia, hypokinesia, manic reaction, myoclonus, psychotic reaction; **Infrequent:** agoraphobia, akathisia, ataxia, CNS depression, convulsion, delirium, delusion, depersonalization, dyskinesia, dystonia, emotional lability, euphoria, extrapyramidal syndrome, gait unsteady, hallucinations, hemiplegia, hostility, hypersomnia, hypochondriasis, hypotonia, hysteria, increased libido, paralysis, paranoid reaction, phobia, psychosis, stupor, twitching, vertigo; **Rare:** akinesia, coma, fibrillations, mutism, obsessions, reflexes decreased, slurred speech, tardive dyskinesia, torticollis, trismus, withdrawal syndrome.

Respiratory System: **Frequent:** cough increased, sinusitis; **Infrequent:** asthma, bronchitis, hoarseness, hyperventilation; **Rare:** apnea, congestion of upper airway, hemoptysis, hiccups, laryngismus, obstructive pulmonary disease, pneumonia.

Skin: **Infrequent:** alopecia, dry skin, exfoliative dermatitis, furunculosis, seborrhea, skin discoloration.

Special Senses: **Infrequent:** accommodation abnormal, conjunctivitis, diplopia, eye pain, mydriasis, otitis media, parosmia, visual field defect; **Rare:** corneal ulcer.

Urogenital System: **Infrequent:** anuria, cystitis, delayed menstruation¹, dysuria, female lactation¹, hematuria, menopause¹, metrorrhagia¹, nocturia, premenstrual syndrome¹, urination impaired, vaginitis¹; **Rare:** kidney calculus, hematospermia², oliguria.

¹ Based on the number of females.

² Based on the number of males.

6.9 Postmarketing Reports

The following adverse reactions have been identified during post-approval use of immediate-release fluvoxamine maleate tablets or LUVOX CR Capsules. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure. (Reactions that are discussed in other sections of this prescribing information are not repeated here.) These reactions include: activation syndrome, aggression, agranulocytosis, anaphylactic reaction, anger, blood glucose increased, bruxism, cardio-respiratory arrest, crying, dysarthria, dysphagia, electrocardiogram QT prolonged, fall, fatigue, feeling drunk, feeling jittery, gait disturbance, gastroesophageal reflux disease, glossodynia, hepatitis, homicidal ideation, impulsive behavior, ileus, inappropriate antidiuretic hormone secretion, interstitial lung disease, irritability, loss of consciousness, lethargy, muscular weakness, Parkinsonism, pancreatitis, pyrexia, renal impairment, rhabdomyolysis, self injurious behavior, shock, somnolence neonatal, Stevens-Johnson syndrome, tachycardia, urinary retention, ventricular arrhythmia, ventricular tachycardia (including torsades de pointes known to cause cardiac arrest, sometimes fatal), vision blurred, white blood cell count decreased.

7 DRUG INTERACTIONS

7.1 Potential Interactions with Drugs that Inhibit or are Metabolized by Cytochrome P450 Isoenzymes

Multiple hepatic cytochrome P450 isoenzymes are involved in the oxidative biotransformation of a large number of structurally different drugs and endogenous compounds. The available knowledge concerning the relationship of fluvoxamine and the cytochrome P450 isoenzyme system has been obtained mostly from pharmacokinetic interaction studies conducted in healthy volunteers, but some preliminary *in vitro* data are also available. Based on a finding of

substantial interactions of fluvoxamine with certain of these drugs (see later parts of this section and also **WARNINGS AND PRECAUTIONS** [5] for details) and limited *in vitro* data for CYP3A4, it appears that fluvoxamine inhibits several cytochrome P450 isoenzymes that are known to be involved in the metabolism of other drugs such as: CYP1A2 (e.g., warfarin, theophylline, propranolol, tizanidine), CYP2C9 (e.g., warfarin), CYP3A4 (e.g., alprazolam), and CYP2C19 (e.g., omeprazole).

In vitro data suggest that fluvoxamine is a relatively weak inhibitor of CYP2D6.

Approximately 7% of the normal population has a genetic code that leads to reduced levels of activity of CYP2D6 enzyme. Such individuals have been referred to as “poor metabolizers” (PM) of drugs such as debrisoquin, dextromethorphan, and tricyclic antidepressants. While none of the drugs studied for drug interactions significantly affected the pharmacokinetics of fluvoxamine, an *in vivo* study of fluvoxamine single-dose pharmacokinetics in 13 PM subjects demonstrated altered pharmacokinetic properties compared to 16 “extensive metabolizers” (EM): mean C_{max} , AUC, and half-life were increased by 52%, 200%, and 62%, respectively, in the PM compared to the EM group. This suggests that fluvoxamine is metabolized, at least in part, by CYP2D6. Caution is indicated in patients known to have reduced levels of cytochrome P450 2D6 activity and those receiving concomitant drugs known to inhibit this cytochrome P450 isoenzyme (e.g., quinidine).

The metabolism of fluvoxamine has not been fully characterized and the effects of potent cytochrome P450 isoenzyme inhibition, such as the ketoconazole inhibition of CYP3A4, on fluvoxamine metabolism have not been studied.

A clinically significant fluvoxamine interaction is possible with drugs having a narrow therapeutic ratio such as pimozide, warfarin, theophylline, certain benzodiazepines, omeprazole, and phenytoin. If LUVOX CR Capsules are to be administered together with a drug that is eliminated via oxidative metabolism and has a narrow therapeutic window, plasma levels and/or pharmacodynamic effects of the latter drug should be monitored closely, at least until steady-state conditions are reached (see **CONTRAINDICATIONS** [4] and **WARNINGS AND PRECAUTIONS** [5]).

7.2 CNS Active Drugs

Antipsychotics: See **WARNINGS AND PRECAUTIONS** (5.3).

Benzodiazepines: See **WARNINGS AND PRECAUTIONS** (5.9).

Alprazolam: See **WARNINGS AND PRECAUTIONS** (5.9).

Diazepam: See **WARNINGS AND PRECAUTIONS** (5.9).

Lorazepam: A study of multiple doses of immediate-release fluvoxamine maleate tablets (50 mg given twice daily) in healthy male volunteers (N=12) and a single dose of lorazepam (4 mg single dose) indicated no significant pharmacokinetic interaction. On average, both lorazepam alone and lorazepam with fluvoxamine produced substantial decrements in cognitive functioning; however, the coadministration of fluvoxamine and lorazepam did not produce larger mean decrements compared to lorazepam alone.

Alcohol: Studies involving single 40 g doses of ethanol (oral administration in one study and intravenous in the other) and multiple dosing with immediate-release fluvoxamine maleate tablets (50 mg given twice daily) revealed no effect of either drug on the pharmacokinetics or pharmacodynamics of the other. As with other psychotropic medications, patients should be advised to avoid alcohol while taking LUVOX CR Capsules.

Carbamazepine: Elevated carbamazepine levels and symptoms of toxicity have been reported with the coadministration of immediate-release fluvoxamine maleate tablets and carbamazepine.

Clozapine: See **WARNINGS AND PRECAUTIONS** (5.9).

Lithium: As with other serotonergic drugs, lithium may enhance the serotonergic effects of fluvoxamine and, therefore, the combination should be used with caution. Seizures have been reported with the coadministration of immediate-release fluvoxamine maleate tablets and lithium.

Methadone: See **WARNINGS AND PRECAUTIONS** (5.9).

Monoamine Oxidase Inhibitors: See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.2).

Pimozide: See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.6).

Ramelteon: See **CONTRAINDICATIONS (4)** and **WARNINGS AND PRECAUTIONS (5.8)**.

Serotonergic Drugs: See **WARNINGS AND PRECAUTIONS (5.3)**.

Tacrine: In a study of 13 healthy, male volunteers, a single 40 mg dose of tacrine added to immediate-release fluvoxamine maleate tablets 100 mg/day administered at steady-state was associated with 5- and 8-fold increases in tacrine C_{max} and AUC, respectively, compared to the administration of tacrine alone. Five subjects experienced nausea, vomiting, sweating, and diarrhea following coadministration, consistent with the cholinergic effects of tacrine.

Thioridazine: See **CONTRAINDICATIONS (4)** and **WARNINGS AND PRECAUTIONS (5.4)**.

Tizanidine: See **CONTRAINDICATIONS (4)** and **WARNINGS AND PRECAUTIONS (5.5)**.

Tricyclic Antidepressants (TCAs): Significantly increased plasma TCA levels have been reported with the coadministration of immediate-release fluvoxamine maleate tablets and amitriptyline, clomipramine or imipramine. Caution is indicated with the coadministration of LUVOX CR Capsules and TCAs; plasma TCA concentrations may need to be monitored, and the dose of TCA may need to be reduced.

Triptans: There have been rare postmarketing reports of serotonin syndrome with use of an SSRI and a triptan. If concomitant treatment of LUVOX CR Capsules with a triptan is clinically warranted, careful observation of the patient is advised, particularly during treatment initiation and dose increases (see **WARNINGS AND PRECAUTIONS (5.3)**).

Sumatriptan - There have been rare postmarketing reports describing patients with weakness, hyperreflexia, and incoordination following the use of a selective serotonin reuptake inhibitor (SSRI) and sumatriptan. If concomitant treatment with sumatriptan and an SSRI (e.g., fluoxetine, fluvoxamine, paroxetine, sertraline) is clinically warranted, appropriate observation of the patient is advised.

Tryptophan: Tryptophan may enhance the serotonergic effects of fluvoxamine, and the combination should, therefore, be used with caution. Severe vomiting has been reported with the coadministration of immediate-release fluvoxamine maleate tablets and tryptophan (see **WARNINGS AND PRECAUTIONS (5.3)**).

7.3 Other Drugs

Alosetron: See **CONTRAINDICATIONS (4)**, **WARNINGS AND PRECAUTIONS (5.7)**, and Lotronex[®] (alosetron) package insert.

Digoxin: Administration of immediate-release fluvoxamine maleate tablets 100 mg daily for 18 days (N=8) did not significantly affect the pharmacokinetics of a 1.25 mg single intravenous dose of digoxin.

Diltiazem: Bradycardia has been reported with the coadministration of immediate-release fluvoxamine maleate tablets and diltiazem.

Mexiletine: See **WARNINGS AND PRECAUTIONS (5.9)**.

Propranolol and Other Beta-Blockers: Coadministration of immediate-release fluvoxamine maleate tablets 100 mg per day and propranolol 160 mg per day in normal volunteers resulted in a mean five-fold increase (range 2 to 17) in minimum propranolol plasma concentrations. In this study, there was a slight potentiation of the propranolol-induced reduction in heart rate and reduction in the exercise diastolic pressure.

One case of bradycardia and hypotension and a second case of orthostatic hypotension have been reported with the coadministration of immediate-release fluvoxamine maleate tablets and metoprolol.

If propranolol or metoprolol is coadministered with LUVOX CR Capsules, a reduction in the initial beta-blocker dose and more cautious dose titration are recommended. No dosage adjustment is required for LUVOX CR Capsules.

Coadministration of immediate-release fluvoxamine maleate tablets 100 mg per day with atenolol 100 mg per day (N=6) did not affect the plasma concentrations of atenolol. Unlike propranolol and metoprolol, which undergo hepatic metabolism, atenolol is eliminated primarily by renal excretion.

Theophylline: See **WARNINGS AND PRECAUTIONS (5.9)**.

Warfarin and Other Drugs That Interfere With Hemostasis (NSAIDs, Aspirin, etc.): See WARNINGS AND PRECAUTIONS (5.9, 5.11).

7.4 Effects of Smoking on Fluvoxamine Metabolism

Smokers had a 25% increase in the metabolism of fluvoxamine compared to nonsmokers.

7.5 Electroconvulsive Therapy (ECT)

There are no clinical studies establishing the benefits or risks of combined use of ECT and fluvoxamine maleate.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Teratogenic Effects – Pregnancy Category C: When pregnant rats were given daily doses of fluvoxamine (60, 120, or 240 mg/kg) orally throughout the period of organogenesis, developmental toxicity in the form of increased embryofetal death and increased incidences of fetal eye abnormalities (folded retinas) was observed at doses of 120 mg/kg or greater. Decreased fetal body weight was seen at the high dose. The no effect dose for developmental toxicity in this study was 60 mg/kg (approximately 2 times the maximum recommended human dose [MRHD] on a mg/m² basis).

In a study in which pregnant rabbits were administered doses of up to 40 mg/kg (approximately 2 times the MRHD on a mg/m² basis) orally during organogenesis, no adverse effects on embryofetal development were observed.

In other reproduction studies in which female rats were dosed orally during pregnancy and lactation (5, 20, 80, or 160 mg/kg), increased pup mortality at birth was seen at doses of 80 mg/kg or greater and decreases in pup body weight and survival were observed at all doses (low effect dose approximately 0.1 times the MRHD on a mg/m² basis).

Nonteratogenic Effects: Neonates exposed to immediate-release fluvoxamine maleate tablets and other SSRIs or serotonin and norepinephrine reuptake inhibitors (SNRIs) late in the third trimester have developed complications requiring prolonged hospitalization, respiratory support, and tube feeding. These findings are based on postmarketing reports. Such complications can arise immediately upon delivery. Reported clinical findings have included respiratory distress, cyanosis, apnea, seizures, temperature instability, feeding difficulty, vomiting, hypoglycemia, hypotonia, hypertonia, hyperreflexia, tremor, jitteriness, irritability and constant crying. These features are consistent with either a direct toxic effect of SSRIs or SNRIs or, possibly, a drug discontinuation syndrome. It should be noted that, in some cases, the clinical picture is consistent with serotonin syndrome (see **WARNINGS AND PRECAUTIONS (5.3)**).

Infants exposed to SSRIs in late pregnancy may have an increased risk for persistent pulmonary hypertension of the newborn (PPHN). PPHN is associated with substantial neonatal morbidity and mortality. In a case-control study of 377 women whose infants were born with PPHN and 836 women whose infants were born healthy, the risk for developing PPHN was approximately 6-fold higher for infants exposed to SSRIs after the 20th week of gestation compared to infants who had not been exposed to antidepressants during pregnancy. PPHN occurs in 1-2 per 1000 live births in the general population.

When treating a pregnant woman with LUVOX CR Capsules during the third trimester, the physician should carefully consider both the potential risks and benefits of treatment (see **DOSAGE AND ADMINISTRATION (2.6)**). Physicians should note that in a prospective longitudinal study of 201 women with a history of major depression who were euthymic at the beginning of pregnancy, women who discontinued antidepressant medication during pregnancy were more likely to experience a relapse of major depression than women who continued antidepressant medication.

8.2 Labor and Delivery

The effect of fluvoxamine on labor and delivery in humans is unknown.

8.3 Nursing Mothers

Fluvoxamine is secreted in human breast milk. Because of the potential for serious adverse reactions in nursing infants from LUVOX CR Capsules, a decision should be made whether to discontinue nursing or discontinue the drug, taking into account the importance of the drug to the mother.

8.4 Pediatric Use

LUVOX CR Capsules have not been evaluated in pediatric patients (see **BOXED WARNING**). The efficacy of fluvoxamine maleate administered as immediate-release tablets for the treatment of OCD was demonstrated in a 10-week multicenter placebo-controlled study with 120 outpatients ages 8-17. In addition, 99 of these outpatients continued open-label fluvoxamine maleate treatment for up to another one to three years, equivalent to 94 patient years. The adverse reaction profile observed in that study was generally similar to that observed in adult studies with immediate-release fluvoxamine maleate tablets (see **ADVERSE REACTIONS (6.3)**).

Decreased appetite and weight loss have been observed in association with the use of fluvoxamine as well as other SSRIs. Consequently, regular monitoring of weight and growth should be performed in children and adolescents treated with an SSRI such as LUVOX CR Capsules.

The risks, if any, that may be associated with fluvoxamine's extended use in children and adolescents with OCD have not been systematically assessed. The prescriber should be mindful that the evidence relied upon to conclude that fluvoxamine is safe for use in children and adolescents derives from relatively short-term clinical studies and from extrapolation of experience gained with adult patients. In particular, there are no studies that directly evaluate the effects of long-term fluvoxamine use on the growth, cognitive behavioral development, and maturation of children and adolescents. Although there is no affirmative finding to suggest that fluvoxamine possesses a capacity to adversely affect growth, development, or maturation, the absence of such findings is not compelling evidence of the absence of the potential of fluvoxamine to have adverse effects in chronic use (see **WARNINGS AND PRECAUTIONS –Clinical Worsening and Suicide Risk [5.1]**).

Safety and effectiveness in the pediatric population other than pediatric patients with OCD have not been established (see **BOXED WARNING** and **WARNINGS AND PRECAUTIONS–Clinical Worsening and Suicide Risk [5.1]**). Anyone considering the use of LUVOX CR Capsules in a child or adolescent must balance the potential risks with the clinical need.

8.5 Geriatric Use

Approximately 230 patients and 5 patients participating in controlled premarketing studies with immediate-release fluvoxamine maleate tablets and LUVOX CR Capsules, respectively, were 65-years of age or over. No overall differences in safety were observed between these patients and younger patients. Other reported clinical experience has not identified differences in response between the elderly and younger patients. However, SSRIs and SNRIs, including fluvoxamine, have been associated with several cases of clinically significant hyponatremia in elderly patients, who may be at greater risk for this adverse reaction (see **WARNINGS AND PRECAUTIONS [5.14]**). Furthermore, the clearance of fluvoxamine is decreased by about 50% in elderly compared to younger patients (see **CLINICAL PHARMACOLOGY–Elderly [12.3]**), and greater sensitivity of some older individuals also cannot be ruled out. Consequently, a lower starting dose should be considered in elderly patients, and LUVOX CR Capsules should be slowly titrated during initiation of therapy.

9 DRUG ABUSE AND DEPENDENCE

9.1 Controlled Substance Class

LUVOX CR is not a controlled substance.

9.2 Physical and Psychological Dependence

The potential for abuse, tolerance, and physical dependence with immediate-release fluvoxamine maleate has been studied in a nonhuman primate model. No evidence of dependency phenomena was found. The discontinuation effects of LUVOX CR Capsules were not systematically evaluated in controlled clinical trials. LUVOX CR Capsules were not systematically studied in clinical trials for potential for abuse, but there was no indication of drug-seeking behavior in clinical trials. It should be noted, however, that patients at risk for drug dependency were systematically excluded from investigational studies of immediate-release fluvoxamine maleate. Generally, it is not possible to predict on the basis of preclinical or premarketing clinical experience the extent to which a CNS active drug will be misused, diverted, and/or abused once marketed. Consequently, health care providers should carefully evaluate patients for a history of drug abuse and follow such patients closely, observing them for signs of LUVOX CR misuse or abuse (i.e., development of tolerance, incrementation of dose, drug-seeking behavior).

10 OVERDOSAGE

10.1 Human Experience

Exposure to immediate-release fluvoxamine maleate tablets includes over 45,000 patients treated in clinical trials and an estimated exposure of 50,000,000 patients treated during worldwide marketing experience (end of 2005). Of the 539 cases of deliberate or accidental overdose involving fluvoxamine reported from this population, there were 55 deaths. Of these, 9 were in patients thought to be taking immediate-release fluvoxamine tablets alone and the remaining 46 were in patients taking fluvoxamine along with other drugs. Among non-fatal overdose cases, 404 patients recovered completely. Five patients experienced adverse sequelae of overdosage, to include persistent mydriasis, unsteady gait, hypoxic encephalopathy, kidney complications (from trauma associated with overdose), bowel infarction requiring a hemicolectomy, and vegetative state. In 13 patients, the outcome was provided as abating at the time of reporting. In the remaining 62 patients, the outcome was unknown. The largest known ingestion of fluvoxamine immediate-release

tablets involved 12,000 mg (equivalent to 2 to 3 months' dosage). The patient fully recovered. However, ingestions as low as 1,400 mg have been associated with lethal outcome, indicating considerable prognostic variability.

In the controlled clinical trials with 403 patients treated with LUVOX CR Capsules, there was one nonfatal intentional overdose.

Commonly ($\geq 5\%$) observed adverse reactions associated with fluvoxamine maleate overdose include gastrointestinal complaints (nausea, vomiting, and diarrhea), coma, hypokalemia, hypotension, respiratory difficulties, somnolence, and tachycardia. Other notable signs and symptoms seen with immediate-release fluvoxamine maleate overdose (single or multiple drugs) include bradycardia, ECG abnormalities (such as heart arrest, QT interval prolongation, first degree atrioventricular block, bundle branch block, and junctional rhythm), convulsions, dizziness, liver function disturbances, tremor, and increased reflexes.

10.2 Management of Overdosage

Treatment should consist of those general measures employed in the management of overdosage with any antidepressant.

Ensure an adequate airway, oxygenation, and ventilation. Monitor cardiac rhythm and vital signs. General supportive and symptomatic measures are also recommended. Induction of emesis is not recommended. Gastric lavage with a large-bore orogastric tube with appropriate airway protection, if needed, may be indicated if performed soon after ingestion, or in symptomatic patients.

Activated charcoal should be administered. Due to the large volume of distribution of this drug, forced diuresis, dialysis, hemoperfusion, and exchange transfusion are unlikely to be of benefit. No specific antidotes for fluvoxamine are known.

A specific caution involves patients taking, or recently having taken, fluvoxamine maleate who might ingest excessive quantities of a tricyclic antidepressant. In such a case, accumulation of the parent tricyclic and/or an active metabolite may increase the possibility of clinically significant sequelae and extend the time needed for close medical observation (see **DRUG INTERACTIONS** [7.2]).

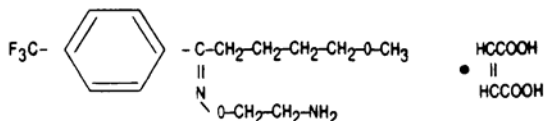
In managing overdosage, consider the possibility of multiple drug involvement. The health care provider should consider contacting a poison control center for additional information on the treatment of any overdose. Telephone numbers for certified poison control centers are listed in the *Physicians' Desk Reference* (PDR).

11 DESCRIPTION

LUVOX CR is an extended-release capsule for oral administration that contains fluvoxamine maleate, a selective serotonin (5-HT) reuptake inhibitor (SSRI) belonging to the chemical series, the 2-aminoethyl oxime ethers of aralkylketones.

Fluvoxamine maleate is chemically designated as 5-methoxy-4'-(trifluoromethyl) valerophenone-(E)-O-(2-aminoethyl)oxime maleate (1:1) and has the empirical formula $C_{15}H_{21}O_2N_2F_3 \cdot C_4H_4O_4$. Its molecular weight is 434.41.

The structural formula is:



Fluvoxamine maleate is a white to off-white, odorless, crystalline powder that is sparingly soluble in water, freely soluble in ethanol and chloroform, and practically insoluble in diethyl ether.

LUVOX CR Capsules are available in 100 mg and 150 mg strengths for oral administration. In addition to the active ingredient, fluvoxamine maleate, each capsule contains the following inactive ingredients: talc, sugar spheres, ammonio methacrylate copolymer type B, dibutyl sebacate, red iron oxide, FD&C Blue No. 2, titanium dioxide, gelatin (porcine- or bovine-derived), and Opacode Grey. LUVOX CR Capsules are gluten-free.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

The mechanism of action of fluvoxamine maleate in obsessive compulsive disorder is presumed to be linked to its specific serotonin reuptake inhibition in brain neurons. Fluvoxamine has been shown to be a potent inhibitor of the serotonin reuptake transporter in preclinical studies, both *in vitro* and *in vivo*.

12.2 Pharmacodynamics

In *in vitro* studies, fluvoxamine maleate had no significant affinity for histaminergic, alpha or beta adrenergic, muscarinic, or dopaminergic receptors. Antagonism of some of these receptors is thought to be associated with various sedative, cardiovascular, anticholinergic, and extrapyramidal effects of some psychotropic drugs.

12.3 Pharmacokinetics

Bioavailability: A single-dose crossover study in 28 healthy subjects was conducted to compare the pharmacokinetics of fluvoxamine after administration of LUVOX CR Capsules and immediate-release fluvoxamine maleate tablets.

In the single-dose crossover study, mean C_{max} was 38% lower and relative bioavailability was 84% for LUVOX CR Capsules versus immediate-release fluvoxamine maleate tablets.

In a multiple-dose proportionality study, LUVOX CR Capsules were administered over a dose range of 100 mg/day to 300 mg/day to 20 healthy volunteers. Steady-state plasma concentrations were achieved within a week of dosing. Mean maximum plasma concentrations were 47 ng/mL, 161 ng/mL, and 319 ng/mL, respectively, at the 100 mg, 200 mg, and 300 mg administered dose levels. Fluvoxamine exhibited nonlinear pharmacokinetics producing disproportionately higher concentrations over the dose range. The AUC and C_{max} values increased 5.7-fold following the 3-fold increase in dose from 100 mg to 300 mg.

Food caused the mean AUC and C_{max} of fluvoxamine to increase only slightly; therefore, administration of LUVOX CR Capsules with food does not significantly affect the absorption of fluvoxamine.

Distribution/Protein Binding: The mean apparent volume of distribution for fluvoxamine is approximately 25 L/kg, suggesting extensive tissue distribution.

Approximately 80% of fluvoxamine is bound to plasma protein, mostly albumin, over a concentration range of 20 to 2000 ng/mL.

Metabolism: Fluvoxamine maleate is extensively metabolized by the liver; the main metabolic routes are oxidative demethylation and deamination. Nine metabolites were identified following a 5 mg radiolabelled dose of fluvoxamine maleate, constituting approximately 85% of the urinary excretion products of fluvoxamine. The main human metabolite was fluvoxamine acid which, together with its N-acetylated analog, accounted for about 60% of the urinary excretion products. A third metabolite, fluvoxethanol, formed by oxidative deamination, accounted for about 10%. Fluvoxamine acid and fluvoxethanol were tested in an *in vitro* assay of serotonin and norepinephrine reuptake inhibition in rats; they were inactive except for a weak effect of the former metabolite on inhibition of serotonin uptake (1-2 orders of magnitude less potent than the parent compound). Approximately 2% of fluvoxamine was excreted in urine unchanged (see **DRUG INTERACTIONS [7]**).

Elimination: Following a ^{14}C -labelled oral dose of fluvoxamine maleate (5 mg), an average of 94% of drug-related products was recovered in the urine within 71 hours.

After administration of a 100 mg, single oral dose of LUVOX CR Capsules, the mean plasma half-life of fluvoxamine in healthy male and female volunteers was 16.3 hours.

Gender: In a study with 15 male and 13 female healthy volunteers who were administered LUVOX CR Capsules 100 mg, AUC and C_{max} of fluvoxamine were increased by approximately 60% in females compared to males. There were no differences in the elimination half-life between males and females.

Elderly Subjects: In a study using immediate-release fluvoxamine maleate tablets at 50 mg and 100 mg and comparing elderly (ages 66-73 years) and young subjects (ages 19-35 years), mean maximum plasma concentrations in the elderly were 40% higher. The multiple-dose elimination half-life of fluvoxamine was 17.4 hours and 25.9 hours in the elderly compared to 13.6 hours and 15.6 hours in the young subjects at steady state for 50 mg and 100 mg doses, respectively.

In elderly patients administered immediate-release fluvoxamine maleate tablets, the clearance of fluvoxamine was reduced by about 50%; therefore, LUVOX CR Capsules should be slowly titrated during initiation of therapy (see **DOSAGE AND ADMINISTRATION [2.3]**).

Pediatric Subjects: The pharmacokinetics of LUVOX CR Capsules have not been evaluated in pediatric patients. However, the multiple-dose pharmacokinetics of fluvoxamine were determined in male and female children (ages 6-11) and adolescents (ages 12-17). Steady-state plasma fluvoxamine concentrations were 2- to 3-fold higher in

children than in adolescents. AUC and C_{max} in children were 1.5- to 2.7-fold higher than that in adolescents (see Table 4). As in adults, both children and adolescents exhibited nonlinear multiple-dose pharmacokinetics. Female children showed significantly higher AUC (0-12) and C_{max} compared to male children and, therefore, lower doses of immediate-release fluvoxamine maleate tablets may produce therapeutic benefit (see Table 5). No gender differences were observed in adolescents. Steady-state plasma fluvoxamine concentrations were similar in adults and adolescents at a dose of 300 mg/day, indicating that fluvoxamine exposure was similar in these two populations (see Table 4). Dose adjustment in adolescents (up to the adult maximum dose of 300 mg) may be indicated to achieve therapeutic benefit.

TABLE 4
COMPARISON OF MEAN (SD) IMMEDIATE-RELEASE TABLET FLUVOXAMINE MALEATE PHARMACOKINETIC PARAMETERS BETWEEN CHILDREN, ADOLESCENTS, AND ADULTS

Pharmacokinetic Parameter (body weight corrected)	Dose = 200 mg/day (100 mg Twice Daily)		Dose = 300 mg/day (150 mg Twice Daily)	
	Children (n = 10)	Adolescent (n = 17)	Adolescent (n = 13)	Adult (n = 16)
AUC 0-12 (ng•h/mL/kg)	155.1 (160.9)	43.9 (27.9)	69.6 (46.6)	59.4 (40.9)
C_{max} (ng/mL/kg)	14.8 (14.9)	4.2 (2.6)	6.7 (4.2)	5.7 (3.9)
C_{min} (ng/mL/kg)	11.0 (11.9)	2.9 (2.0)	4.8 (3.8)	4.6 (3.2)

TABLE 5
COMPARISON OF MEAN (SD) IMMEDIATE-RELEASE TABLET FLUVOXAMINE MALEATE PHARMACOKINETIC PARAMETERS BETWEEN MALE AND FEMALE CHILDREN (6-11 YEARS)

Pharmacokinetic Parameter (body weight corrected)	Dose = 200 mg/day (100 mg Twice Daily)	
	Male Children (n = 7)	Female Children (n = 3)
AUC 0-12 (ng•h/mL/kg)	95.8 (83.9)	293.5 (233.0)
C_{max} (ng/mL/kg)	9.1 (7.6)	28.1 (21.1)
C_{min} (ng/mL/kg)	6.6 (6.1)	21.2 (17.6)

Hepatic and Renal Disease: A cross-study comparison (healthy subjects versus patients with hepatic dysfunction) using immediate-release fluvoxamine maleate tablets suggested a 30% decrease in fluvoxamine clearance in association with hepatic dysfunction. The mean minimum plasma concentrations in renally impaired patients (creatinine clearance of 5 mL/min to 45 mL/min) after 4 weeks and 6 weeks of treatment (50 mg given twice daily, N = 13) were comparable to each other, suggesting no accumulation of fluvoxamine in these patients (see **WARNINGS AND PRECAUTIONS–Use in Patients with Concomitant Illness [5.15]**).

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenesis: There was no evidence of carcinogenicity in rats treated orally with fluvoxamine maleate for 30 months or hamsters treated orally with fluvoxamine maleate for 20 months (females) or 26 months (males). The daily doses in the high-dose groups in these studies were increased over the course of the study from a minimum of 160 mg/kg to a maximum of 240 mg/kg in rats, and from a minimum of 135 mg/kg to a maximum of 240 mg/kg in hamsters. The maximum dose of 240 mg/kg is approximately 6 times the maximum recommended human dose (MRHD) on a mg/m² basis.

Mutagenesis: No evidence of genotoxic potential was observed in a mouse micronucleus test, an *in vitro* chromosome aberration test, or the Ames microbial mutagen test with or without metabolic activation.

Impairment of Fertility: In a study in which male and female rats were administered fluvoxamine (60, 120, or 240 mg/kg) orally prior to and during mating and gestation, fertility was impaired at doses of 120 mg/kg or greater, as

evidenced by increased latency to mating, decreased sperm count, decreased epididymal weight, and decreased pregnancy rate. In addition, the numbers of implantations and embryos were decreased at the highest dose. The no effect dose for fertility impairment was 60 mg/kg (approximately 2 times the MRHD on a mg/m² basis).

14 CLINICAL STUDIES

14.1 Obsessive Compulsive Disorder (OCD)

The effectiveness of LUVOX CR Capsules for the treatment of OCD was demonstrated in a 12-week, multicenter, placebo-controlled study of adult outpatients. Patients in this trial were titrated in 50 mg increments over the first six weeks of the study on the basis of response and tolerance from a dose of 100 mg/day to a fluvoxamine maleate dose within a range of 100 mg to 300 mg once-a-day. Patients in this study had moderate to severe OCD (DSM-IV), with mean baseline ratings on the Yale-Brown Obsessive Compulsive Scale (Y-BOCS), total scores of 26.6 and 26.3 for fluvoxamine and placebo-treatment groups, respectively.

Patients receiving LUVOX CR Capsules demonstrated statistically significant improvement over placebo patients at the primary endpoint (Week 12) compared to baseline on the Y-BOCS. The mean daily dose of LUVOX CR Capsules administered to patients was 261 mg at end of study.

Exploratory analyses for age and gender effects on outcomes did not show any significant differential responsiveness on the basis of age or sex.

The effectiveness of immediate-release fluvoxamine maleate tablets for the treatment of OCD was demonstrated in two 10-week multicenter, parallel group studies of adult outpatients. Patients in these trials were titrated to a total daily fluvoxamine maleate dose of 150 mg/day over the first two weeks of the trial, after which the dose was adjusted within a range of 100-300 mg/day (given in two doses per day), on the basis of response and tolerance. Patients in these studies had moderate to severe OCD (DSM-III-R), with mean baseline ratings on the Yale-Brown Obsessive Compulsive Scale (Y-BOCS) total score of 23.

14.2 Adult OCD Maintenance Study with Immediate-Release Fluvoxamine Maleate Tablets

In a maintenance trial of adult outpatients with OCD, 114 patients meeting DSM-IV criteria for OCD and with a Yale-Brown Obsessive Compulsive Scale (Y-BOCS) score ≥ 18 were titrated to an effective dose of immediate-release fluvoxamine maleate tablets 100 to 300 mg/day as part of an initial 10-week single-blind treatment phase. Treatment response during this single-blind phase was defined as Y-BOCS scores at least 30% lower than baseline at the end of weeks 8 and 10. Of the patients who responded, their average duration of response was 4 weeks. Patients who responded during this initial phase were randomized either to continuation of immediate-release fluvoxamine maleate tablets (N=56) or to placebo (N=58) in a double-blind phase for observation of relapse. Relapse during the double-blind phase was defined as an increase in the Y-BOCS score of at least 30% over the baseline for that phase or patient refusal to continue treatment due to a substantial increase in OCD symptoms. In the double-blind phase, patients receiving continued immediate-release fluvoxamine maleate tablets treatment experienced, on average, a significantly lower relapse rate than those receiving placebo.

An examination of population subgroups from this trial did not reveal any clear evidence of a differential maintenance effect on the basis of age or gender.

14.3 Pediatric OCD Study

LUVOX CR Capsules have not been evaluated in pediatric patients. However, the effectiveness of immediate-release fluvoxamine maleate tablets for the treatment of OCD was demonstrated in a 10-week multicenter, parallel group study in a pediatric outpatient population (children and adolescents, ages 8-17 years). Patients in this study were titrated to a total daily fluvoxamine dose of approximately 100 mg/day over the first two weeks of the trial, following which the dose was adjusted within a range of 50-200 mg/day (given in two doses per day) on the basis of response and tolerance. All patients had moderate-to-severe OCD (DSM-III-R) with mean baseline ratings on the Children's Yale-Brown Obsessive Compulsive Scale (CY-BOCS) total score of 24.

Post hoc exploratory analyses for gender effects on outcomes did not suggest any differential responsiveness on the basis of gender. Further exploratory analyses revealed a prominent treatment effect in the 8 year to 11 year age group and essentially no effect in the 12 year to 17 year age group. While the significance of these results is not clear, the 2-3 fold higher steady-state plasma fluvoxamine concentrations in children compared to adolescents (see *CLINICAL PHARMACOLOGY-Pediatric Subjects [12.3]*) is suggestive that decreased exposure in adolescents may have been a factor, and dose adjustment in adolescents (up to the adult maximum dose of 300 mg/day) may be indicated to achieve therapeutic benefit.

16 HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied

LUVOX CR Capsules are available in the following strengths, colors, imprints, and presentations:

100 mg Extended-Release Capsules: Available in a two-piece gelatin capsule (dark blue opaque cap/white opaque body) imprinted with $\text{\textcircled{P}}$ on one side of the cap and LCR 100 on the other side of the cap.

Bottles of 30.....NDC 68727-600-01

150 mg Extended-Release Capsules: Available in a two-piece gelatin capsule (dark blue opaque cap/powder blue opaque body) imprinted with $\text{\textcircled{P}}$ on one side of the cap and LCR 150 on the other side of the cap.

Bottles of 30.....NDC 68727-601-01

16.2 Storage

Keep out of reach of children.

LUVOX CR Capsules should be protected from high humidity and stored at 25°C (77°F); excursions permitted to 15°-30°C (59°-86°F) [see USP Controlled Room Temperature].

Avoid exposure to temperatures above 30°C (86°F).

Dispense in tight containers.

17 PATIENT COUNSELING INFORMATION

See FDA-approved *Medication Guide* (17.10).

Prescribers or other health professionals should inform patients, their families, and their caregivers about the benefits and risks associated with treatment with LUVOX CR Capsules and should counsel them in the appropriate use. A patient Medication Guide discussing antidepressant medicines, depression and other serious mental illnesses, and suicidal thoughts or actions and other important information about LUVOX CR Capsules is available for LUVOX CR Capsules. The prescriber or health professional should instruct patients, their families, and their caregivers to read both sections of the Medication Guide and should assist them in understanding its contents. Patients should be given the opportunity to discuss the contents of the Medication Guide and to obtain answers to any questions they may have. The complete text of the Medication Guide is reprinted at the end of this document.

Patients should be advised of the following issues and asked to alert their prescriber if these occur while taking LUVOX CR Capsules.

17.1 Clinical Worsening and Suicide Risk

Patients, their families, and their caregivers should be encouraged to be alert to the emergence of anxiety, agitation, panic attacks, insomnia, irritability, hostility, aggressiveness, impulsivity, akathisia (psychomotor restlessness), hypomania, mania, other unusual changes in behavior, worsening of depression, and suicidal ideation, especially early during antidepressant treatment and when the dose is adjusted up or down. Families and caregivers of patients should be advised to look for the emergence of such symptoms on a day-to-day basis, since changes may be abrupt. Such symptoms should be reported to the patient's prescriber or health professional, especially if they are severe, abrupt in onset, or were not part of the patient's presenting symptoms. Symptoms such as these may be associated with an increased risk for suicidal thinking and behavior and indicate the need for very close monitoring and possibly changes in the medication (see **BOXED WARNING** and **WARNINGS AND PRECAUTIONS** [5.1]).

17.2 Contraindicated Medications

Patients should be advised that the following medications should not be used while taking LUVOX CR Capsules:

- **Monoamine oxidase inhibitors (MAOIs):** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.2).
- **Thioridazine:** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.4).
- **Tizanidine:** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.5).
- **Pimozide:** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.6).
- **Alosetron:** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.7).
- **Ramelteon:** See **CONTRAINDICATIONS** (4) and **WARNINGS AND PRECAUTIONS** (5.8).

In addition, MAOIs should not be taken within 14 days (2 weeks) after stopping LUVOX CR Capsules, and LUVOX CR Capsules should not be taken within two weeks after stopping treatment with an MAOI (see **CONTRAINDICATIONS** [4] and **WARNINGS AND PRECAUTIONS** [5.2]).

17.3 Other Potentially Hazardous Drug Interactions

Patients should be advised that the use of LUVOX CR Capsules with any of the following medications may produce clinically significant adverse reactions. Patients should inform their physician if they are taking any of these medications before starting treatment with LUVOX CR Capsules. Patients should also inform their physician prior to taking any of these medications while receiving LUVOX CR Capsule therapy.

- **Serotonergic drugs, including triptans, tramadol, and tryptophan:** See *WARNINGS AND PRECAUTIONS (5.3)*.
- **Antipsychotic agents, including clozapine:** See *WARNINGS AND PRECAUTIONS (5.3 and 5.9)*.
- **Certain benzodiazepines:** See *WARNINGS AND PRECAUTIONS (5.9)*.
- **Methadone:** See *WARNINGS AND PRECAUTIONS (5.9)*.
- **Mexiletine:** See *WARNINGS AND PRECAUTIONS (5.9)*.
- **Theophylline:** See *WARNINGS AND PRECAUTIONS (5.9)*.
- **Warfarin and other drugs that interfere with hemostasis:** Patients should be cautioned about the concomitant use of fluvoxamine and NSAIDs, aspirin, or other drugs that affect coagulation since the combined use of psychotropic drugs that interfere with serotonin reuptake and these agents has been associated with an increased risk of bleeding (see *WARNINGS AND PRECAUTIONS [5.9 and 5.11]*).
- **Diuretics:** See *WARNINGS AND PRECAUTIONS (5.14)*.

In addition, patients should be advised to notify their physicians if they are taking, or plan to take, any prescription or over-the-counter drugs, since there is a potential for clinically important interactions with LUVOX CR Capsules.

17.4 Abnormal Bleeding

Patients should be advised that LUVOX CR Capsules may increase the risk of bleeding events, which have ranged from ecchymoses, hematomas, epistaxis, and petechiae to life-threatening hemorrhages. Concomitant use of aspirin, nonsteroidal anti-inflammatory drugs, warfarin, and other anticoagulants may add to this risk (see *WARNINGS AND PRECAUTIONS [5.9 and 5.11]*).

17.5 Interference with Cognitive or Motor Performance

Since any psychoactive drug may impair judgment, thinking, or motor skills, patients should be cautioned about operating hazardous machinery, including automobiles, until they are certain that LUVOX CR Capsules therapy does not adversely affect their ability to engage in such activities.

17.6 Pregnancy

Patients should be advised to notify their physicians if they become pregnant or intend to become pregnant during therapy with LUVOX CR Capsules (see *USE IN SPECIFIC POPULATIONS [8.1]*).

17.7 Nursing

Patients receiving LUVOX CR Capsules should be advised to notify their physicians if they are breast-feeding an infant. (See *USE IN SPECIFIC POPULATIONS–Nursing Mothers [8.3]*.)

17.8 Alcohol

As with other psychotropic medications, patients should be advised to avoid alcohol while taking LUVOX CR Capsules.

17.9 Allergic Reactions

Patients should be advised to notify their physicians if they develop a rash, hives, or a related allergic phenomenon during therapy with LUVOX CR Capsules.

Lotronex[®] is a registered trademark of GlaxoSmithKline.

LUVOX CR[®] is a registered trademark of Abbott Products, Inc.

Distributed by:

Jazz Pharmaceuticals, Inc.
Palo Alto, CA 94304

17.10 FDA-Approved Medication Guide

Medication Guide **LUVOX CR® (LOO Vox CR)** **(Fluvoxamine Maleate) Extended-Release Capsules**

Read the Medication Guide that comes with LUVOX CR Capsules before you start taking it and each time you get a refill. There may be new information. This Medication Guide does not take the place of talking to your healthcare provider about your medical condition or treatment. Talk to your healthcare provider if there is something you do not understand or want to learn more about.

What is the most important information I should know about LUVOX CR Capsules?

LUVOX CR is the same kind of medicine as those used to treat depression. These medicines may cause serious side effects, including:

1. Suicidal thoughts or actions:

- **LUVOX CR Capsules and other antidepressant medicines may increase suicidal thoughts or actions** in some children, teenagers, or young adults within the **first few months of treatment or when the dose is changed**.
- Depression or other serious mental illnesses are the most important causes of suicidal thoughts or actions.
- Watch for these changes and call your healthcare provider right away if you notice:
 - New or sudden changes in mood, behavior, actions, thoughts, or feelings, especially if severe.
 - Pay particular attention to such changes when LUVOX CR is started or when the dose is changed.

Keep all follow-up visits with your healthcare provider and call between visits if you are worried about symptoms.

Call your healthcare provider right away if you have any of the following symptoms, or call 911 in an emergency, especially if they are new, worse, or worry you:

- attempts to commit suicide
- acting on dangerous impulses

- acting aggressive or violent
- thoughts about suicide or dying
- new or worse depression
- new or worse anxiety or panic attacks
- feeling agitated, restless, angry or irritable
- trouble sleeping
- an increase in activity or talking more than what is normal for you
- other unusual changes in behavior or mood

Call your healthcare provider right away if you have any of the following symptoms, or call 911 in an emergency. LUVOX CR Capsules may be associated with these serious side effects:

- 2. Serotonin Syndrome or Neuroleptic Malignant Syndrome-like reactions. This condition can be life-threatening and may include:**
 - agitation, hallucinations, coma or other changes in mental status
 - coordination problems or muscle twitching (overactive reflexes)
 - racing heartbeat, high or low blood pressure
 - sweating or fever
 - nausea, vomiting, or diarrhea
 - muscle rigidity
- 3. Severe allergic reactions:**
 - trouble breathing
 - swelling of the face, tongue, eyes, or mouth
 - rash, itchy welts (hives) or blisters, alone or with fever or joint pain
- 4. Abnormal bleeding:** LUVOX CR Capsules and other antidepressant medicines may increase your risk of bleeding or bruising, especially if you take the blood thinner warfarin (Coumadin®),

Jantoven[®]), a non-steroidal anti-inflammatory drug (NSAIDs, like ibuprofen or naproxen), or aspirin.

5. Seizures or convulsions

6. Manic episodes:

- greatly increased energy
- severe trouble sleeping
- racing thoughts
- reckless behavior
- unusually grand ideas
- excessive happiness or irritability
- talking more or faster than usual

7. Changes in appetite or weight.

Children and adolescents should have height and weight monitored during treatment.

8. Low salt (sodium) levels in the blood.

Elderly people may be at greater risk for this. Symptoms may include:

- headache
- weakness or feeling unsteady
- confusion, problems concentrating or thinking, or memory problems

Do not stop taking LUVOX CR Capsules without first talking to your healthcare provider. Stopping LUVOX CR Capsules may cause serious symptoms, including:

- anxiety, irritability, high or low mood, feeling restless or changes in sleep habits
- headache, sweating, nausea, dizziness
- electric shock-like sensations, shaking, confusion

What is LUVOX CR Capsules?

LUVOX CR Capsules is a prescription medicine used to treat obsessive compulsive disorder (OCD). It is the same kind of drug that is used to treat depression. It is important to talk with your healthcare provider about the risks of treating OCD and also the risks of not treating it. You should discuss all treatment choices with your healthcare provider.

Talk to your healthcare provider if you do not think that your condition is getting better with LUVOX CR treatment.

Who should not take LUVOX CR Capsules?

Do not take LUVOX CR Capsules if you:

- are allergic to fluvoxamine maleate or any of the ingredients in LUVOX CR Capsules. See the end of this Medication Guide for a complete list of ingredients in LUVOX CR Capsules.
- take a Monoamine Oxidase Inhibitor (MAOI). Ask your healthcare provider or pharmacist if

you are not sure if you take an MAOI, including the antibiotic linezolid.

- Do not take an MAOI within 2 weeks of stopping LUVOX CR Capsules.
- Do not start LUVOX CR Capsules if you stopped taking an MAOI in the last 14 days (2 weeks).

People who take LUVOX CR Capsules close in time to an MAOI may have serious or even life-threatening side effects. Get medical help right away if you have any of these symptoms:

- high fever
- uncontrolled muscle spasms
- stiff muscles
- rapid changes in heart rate or blood pressure
- confusion
- loss of consciousness (pass out)
- **take Mellaril[®] (thioridazine). Mellaril[®] should not be taken with LUVOX CR Capsules because this can cause serious heart rhythm problems or sudden death.**
- **take Zanaflex[®] (tizanidine) because LUVOX CR can increase the amount of Zanaflex in your body, which could increase its actions and side effects. This could include causing drowsiness and a drop in blood pressure and affecting how well you do things that require alertness.**
- **take the antipsychotic medicine Orap[®] (pimozide) because this can cause serious heart problems.**
- **Take Lotronex[®] (alosetron) because LUVOX CR can increase the amount of Lotronex in your body, which could increase its actions and side effects.**
- **take Rozerem[®] (ramelteon) because LUVOX CR can increase the amount of Rozerem in your body, which could increase its actions and side effects.**

What should I tell my healthcare provider before taking LUVOX CR Capsules? Ask if you are not sure.

Before starting LUVOX CR Capsules, tell your healthcare provider if you:

- are taking certain drugs such as:
 - Clozaril[®] (clozapine): used to treat schizophrenia
 - Mexitil[®] (mexiletine): used to treat problems with heart rhythm
 - Triptans: used to treat migraine headache
 - Medicines used to treat mood, anxiety, psychotic or thought disorders, including

tricyclics, lithium, SSRIs, SNRIs, or antipsychotics

- Tramadol: used to reduce pain
- Benzodiazepines: used to reduce anxiety, stress, emotional upset, or seizures; helps you sleep; helps with alcohol withdrawal; reduces restlessness; and relaxes muscles
- Methadone: used to relieve pain or to help with addiction
- Theophylline used to treat swollen air passages in your lungs, to relax the muscles in your chest to ease shortness of breath, often to treat asthma
- Warfarin and other drugs that affect how your blood clots
- Diuretics to treat high blood pressure, congestive heart failure, or swelling
- Over-the-counter supplements such as tryptophan or St. John's Wort
- have liver problems
- have kidney problems
- have heart problems
- have or had seizures or convulsions
- have bipolar disorder or mania
- have low sodium levels in your blood
- have a history of a stroke
- have high blood pressure
- have or had bleeding problems
- are pregnant or plan to become pregnant. It is not known if LUVOX CR Capsules will harm your unborn baby. Talk to your healthcare provider about the benefits and risks of treating obsessive compulsive disorder (OCD) during pregnancy
- are breast-feeding or plan to breast-feed. Some LUVOX CR may pass into your breast milk. Talk to your healthcare provider about the best way to feed your baby while taking LUVOX CR Capsules.

Tell your healthcare provider about all the medicines that you take, including prescription and non-prescription medicines, vitamins, and herbal supplements. LUVOX CR Capsules and some medicines may interact with each other, may not work as well, or may cause serious side effects.

Your healthcare provider or pharmacist can tell you if it is safe to take LUVOX CR Capsules with your other medicines. Do not start or stop any medicine while taking LUVOX CR Capsules without talking to your healthcare provider first.

If you take LUVOX CR Capsules, you should not take any other medicines that contain fluvoxamine maleate including: Fluvoxamine Maleate Immediate-Release Tablets.

How should I take LUVOX CR Capsules?

- Take LUVOX CR Capsules at night exactly as prescribed. Your healthcare provider may need to change the dose of LUVOX CR Capsules until it is the right dose for you.
- LUVOX CR Capsules may be taken with or without food.
- If you miss a dose of LUVOX CR Capsules, take the missed dose as soon as you remember. If it is almost time for the next dose, skip the missed dose and take your next dose at the regular time. Do not take two doses of LUVOX CR Capsules at the same time.
- If you take too much LUVOX CR, call your healthcare provider or poison control center right away, or get emergency treatment.

What should I avoid while taking LUVOX CR Capsules?

LUVOX CR Capsules can cause sleepiness or may affect your ability to make decisions, think clearly, or react quickly. You should not drive, operate heavy machinery, or do other dangerous activities until you know how LUVOX CR Capsules affect you. Do not drink alcohol while using LUVOX CR Capsules.

What are the possible side effects of LUVOX CR Capsules?

LUVOX CR may cause serious side effects, including all of those described in the section entitled "What is the most important information I should know about LUVOX CR?"

Common possible side effects in people who take fluvoxamine include:

- Nausea
- Sleepiness
- Weakness
- Dizziness
- Feeling anxious
- Trouble sleeping
- Sexual problems
- Sweating
- Shaking
- Not feeling hungry
- Dry mouth
- Diarrhea
- Muscle pain
- Sore throat

- Throwing up
- Upset stomach
- Yawning

Other side effects in children and adolescents taking fluvoxamine include:

- abnormal increase in muscle movement or agitation
- depression
- heavy menstrual periods
- flatulence (gas)
- rash

Tell your healthcare provider if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of LUVOX CR Capsules. For more information, ask your healthcare provider or pharmacist.

CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT SIDE EFFECTS. YOU MAY REPORT SIDE EFFECTS TO FDA AT 1-800-FDA-1088.

How should I store LUVOX CR Capsules?

- Store LUVOX CR Capsules at 59°F to 86°F (15°C to 30°C).
- Keep LUVOX CR Capsules away from high temperatures (above 86°F or 30°C) and high humidity (dampness).
- Keep the LUVOX CR Capsules bottle closed tightly.

Keep LUVOX CR Capsules and all medicines out of the reach of children.

General information about LUVOX CR Capsules

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use LUVOX CR Capsules for a condition for which it was not prescribed. Do not give LUVOX CR Capsules to other people, even if they have the same condition. It may harm them.

This Medication Guide summarizes the most important information about LUVOX CR Capsules. If you would like more information, talk with your healthcare provider. You may ask your healthcare provider or pharmacist for information about LUVOX CR Capsules that is written for healthcare professionals.

For more information about LUVOX CR Capsules, call (1-800-520-5568) or go to jazzsafety@jazzpharma.com.

What are the ingredients in LUVOX CR Capsules?

Active ingredient: fluvoxamine maleate

Inactive ingredients:

- **Extended-Release Capsules:** ammonio methacrylate copolymer type B, dibutyl sebacate, colorants (FD&C Blue No. 2, Opacode Grey, red iron oxide, titanium dioxide), gelatin (porcine- or bovine-derived), sugar spheres, and talc. LUVOX CR Capsules are gluten-free.

Distributed by Jazz Pharmaceuticals, Palo Alto CA 94304

LCR-MG Rev 0511

This Medication Guide has been approved by the U.S. Food and Drug Administration.

LUVOX CR[®] is a registered trademark of Abbott Products, Inc.